EXPERIMENTAL STUDY

Regulation of ghrelin secretion by somatostatin analogs in rats

Antonio P Silva, Kerstin Bethmann, Friedrich Raulf and Herbert A Schmid
Novartis Institutes for Biomedical Research, Basel, Switzerland
(Correspondence should be addressed to H Schmid; Email: herbert.schmid@novartis.com)

Abstract

Objective: Ghrelin is a hormone present in the plasma in two forms: octanoylated and des-octanoylated ghrelin. In pathophysiological conditions such as Prader–Willi syndrome and ghrelinoma, elevated ghrelin plasma levels are associated with pathological obesity. Clinical studies have shown that somatostatin downregulates ghrelin plasma levels in healthy volunteers. The aim of this study was to investigate the effects of two somatostatin analogues, SOM230 and octreotide, on ghrelin secretion in rats.

Methods: Ghrelin secretion was either unstimulated or stimulated by overnight fasting. Treatment with SOM230 and octreotide was either acute (s.c. injection 1 h before blood sampling) or prolonged (continuous s.c. infusion via 14-day osmotic minipumps).

Results: Acute treatment with octreotide dose-dependently inhibited unstimulated and stimulated secretion of total and active ghrelin. SOM230 (30 μg/kg) inhibited active ghrelin in fasted rats. Lower doses had no effect. After 7 days of treatment, active ghrelin was strongly inhibited by both compounds in fasted animals, with a stronger effect for octreotide. Lower inhibition was achieved in fed rats. After 14 days, the inhibition with octreotide in fasted rats was lower and SOM230 had no effect. Somatostatin receptor expression analysis in the rat glandular stomach revealed a predominant sst₁ and sst₂ expression, low expression of sst₃ and sst₄, and hardly detectable sst₅ mRNA expression.

Conclusions: Somatostatin analogues may be useful for the inhibition of physiologically elevated ghrelin plasma levels. This inhibition appears to be mediated by sst₂ receptors in the rat, and desensitizes after 14 days of treatment.

European Journal of Endocrinology 152 887–894

Introduction

Ghrelin is a 28-amino-acid peptide identified in 1999 and produced mainly in the stomach (1). It is the endogenous ligand for the growth hormone secretagogue receptors (GHS-Rs). Beside its growth hormone (GH)-releasing activity, ghrelin exerts additional effects, such as increased food intake and adiposity, stimulation of stomach motility and secretion, inhibition of insulin secretion, and cardioprotective effects. These effects are mediated by the receptor GHS-R1a, and exerted by acylated ghrelin (also called active ghrelin, octanoylated on Ser³) (2). Des-octanoyl-ghrelin also has biological activity; indeed, antiproliferative effects on the human breast carcinoma cell line MCF-7 have been reported (3). The regulation of energy balance by ghrelin is controlled by the modulation of ghrelin plasma levels, which are upregulated by fasting, hypoglycemia, and leptin, and downregulated by food intake, hyperglycemia, and obesity (4–9).

In pathophysiological conditions, such as Prader–Willi syndrome (PWS) and ghrelinoma, elevated plasma levels of ghrelin are reported (10–13). PWS is a genetic disorder affecting 1 in 15 000 children. The main features of the syndrome are infantile hypotonia, mental deficiency, hypogonadism, and obesity associated with uncontrollable and voracious appetite (14). Whether the high levels of ghrelin reported in these patients are responsible for the obese phenotype is not established clearly. However, the orexigenic effect of ghrelin has been clearly established in humans (15, 16). Recent publications have shown the production of ghrelin in endocrine tumors of the gastrointestinal tract (13, 17, 18). In these studies, the elevated plasma levels of ghrelin observed in patients presenting with ghrelinoma coincided with a high body mass index (≥ 30 kg/m²). Ghrelin secretion can be regulated by different mediators including leptin, estrogens, insulin, glucose, acetylcholine, and somatostatin (2, 5, 19). In human
healthy volunteers, somatostatin was shown to downregulate elevated plasma levels of ghrelin in subjects fasted for 12 or 36 h (20, 21). Somatostatin is a 14- or 28-amino-acid peptide involved in the regulation of hormone and growth factor secretion in various organs. Currently available somatostatin analogues (octreotide and lanreotide) are used for the treatment of two major indications, i.e. acromegaly and carcinoid syndrome, associated with gastro-entero-pancreatic tumors. Since some 50% of gastro-entero-pancreatic tumor patients show an escape of response after 1 year of treatment and 35% of acromegalic patients do not or only partially respond to treatment with these sst2-prefering agonists (22, 23), a new somatostatin analogue with a wider affinity profile for sst receptors was developed. SOM230 is a cyclohexapeptide able to bind sst1, sst2, sst3, and sst5 receptors with high affinity (24). This novel multiligand compound shows a 30- and 40-fold higher affinity to sst1 and sst5, respectively, and has a 3-fold lower affinity to sst2 compared with octreotide. SOM230 is very effective in lowering GH secretion from the pituitary in vitro as well as in vivo. Insulin-like growth factor I (IGF-I) secretion is also decreased strongly by SOM230 in rats (25). Due to its prolonged terminal elimination half-life (23 h in rats, 11 h in humans), SOM230 has a very long duration of action compared with octreotide (25). Furthermore, the escape of response observed for the control of IGF-I secretion on treatment with octreotide is not observed with SOM230 in rats (26). The aim of this study was to determine the comparative effects of octreotide and SOM230 on the secretion of both total ghrelin and active ghrelin in rats either fasted overnight or fed ad libitum.

Materials and methods

The experiments described in this article were performed according to national animal welfare requirements (Eidgenossische Tierversuchsbewilligung, Switzerland).

Compounds and formulation

For acute (1 h) treatment by s.c. injection, SOM230 acetate was dissolved in sterile water and stocks stored frozen in 100× concentrations. On the day of experiment, the stock solutions were diluted with 0.9% sterile saline to the final concentration (pH 5–6). SMS 201–995 acetate (octreotide) was prepared in the same way as SOM230. For the long-term (14 days) treatment, SOM230 and octreotide were dissolved in sterile saline and put in Alzet osmotic 14-day minipumps delivering SOM230 or octreotide at a dose of 3, 10, or 30 μg/kg per h.

Animals and treatments

Adult male Lewis rats (LEW/Han/Hsd, Harlan, The Netherlands), 236–298 g, were housed three per cage under controlled light (12 h light/12 h dark) and temperature (22°C). The rats had water and chow food ad libitum unless otherwise stated. Animals were weighed on the day of blood sampling, and on the day of minipump insertion for the long-term-treatment experiments. For the 1-h treatment experiment all animals, except those of the fed groups, were fasted at 1500 h on the day before the experiment. On the day of experiment, SOM230 or octreotide was injected s.c., 1 ml/kg, at a dose of 3, 10, or 30 μg/kg. After 1 h, animals were anesthetized with 5% isoflurane (Forene; Abbott AG, Baar, Switzerland), and total blood was taken from the abdominal aorta. For the 14-day treatment experiment, minipumps containing either 0.9% NaCl, SOM230 or octreotide were inserted s.c. on day 0. On days 6 and 13, animals (except those in the fed groups) were fasted at 1500 h. On day 7, after anesthesia with 5% isoflurane, total blood was taken from the sublingual vein. On day 14, after anesthesia with 5% isoflurane, total blood was taken from the abdominal aorta. For ghrelin (total and active), plasma was taken on EDTA and aprotinin (100 U/ml). Plasma was separated by centrifugation at 3000 r.p.m. for 10 min at 4°C, and aliquoted in different tubes for total and active ghrelin. For the determination of active ghrelin, plasma was acidified with 10 μl HCl (1 M)/100 μl plasma to avoid hydrolysis of the octanoyl group of active ghrelin.

Determination of total and active ghrelin

Plasma levels of total and active ghrelin were determined by RIA, using commercially available kits (Linco Research, St Charles, MI, USA; catalogue nos GHRT89HK and GHRA88HK, respectively).

Somatostatin receptor expression analysis

Lewis rats (LEW/Han/Hsd; five per group) were either fed ad libitum or fasted overnight. On the day of experiment, they were anesthetized with 5% isoflurane, and killed with an overdose of pentobarbital (Vetanarcol; Veterinaria AG, Zurich, Switzerland). The stomach was excised, opened longitudinally, and a biopsy of the glandular part of the stomach was snap-frozen in liquid nitrogen and stored at −80°C. Frozen samples were incubated for at least 18 h at −20°C in RNALater-ICE (Ambion, Austin, TX, USA). Tissue disruption and homogenization was performed in guanidine thiocyanate containing lysis buffer (RT buffer, RNasy minikit; Qiagen, Basel, Switzerland) using a BioPulverizer Green tube (BIO 101, Carlsbad, CA, USA). Total RNA was then extracted using the RNasy minikit according to manufacturer’s instructions, including a
digestion with DNase to minimize contamination with genomic DNA. 1 μg total RNA was reverse transcribed using the superscript III reverse transcriptase (Innitrogen, Basel, Switzerland) and random hexamer primers (Roche Diagnostics, Basel, Switzerland). The resulting cDNA (amount equivalent to 10 ng total RNA) was submitted to quantitative real-time PCR for the following targets: sst1, sst2, sst3, sst4, sst5, and 18 S rRNA. The primers and probes for somatostatin and sst2 were provided as TaqMan assays-on-demand (Applied Biosystems, Foster City, CA, USA; catalogue nos Rn_00561967_m1 and Rn_00571116_m1, respectively). Primers and probe for sst1, sst3, sst4, and sst5 were provided as assays-by-design (Applied Biosystems).

The sequences for TaqMan primers and probes were as follows. sst1, forward primer, 5'-AGA GCT GGA ACC AGA CTG T-3'; reverse primer, 5'-CAG TCA GCA GTA ACT AG-3'; probe, 5'-AGA GAT CAG TCA GCA GAT AGG TGG-3'; sst2, forward primer, 5'-GCC CGG TAG GT-3'; reverse primer, 5'-CCC TCT AGG AAC-3'; probe, 5'-GCC CGG TAG GT-3'; sst3, forward primer, 5'-GGT TCG TCT CAC CTT-3'; reverse primer, 5'-GAG TGG CTC TGA TCT-3'; probe, 5'-GAG TGG CTC TGA TCT-3'; sst4, forward primer, 5'-GCC TCT CCT CTC GGA CAA CT-3'; reverse primer, 5'-GGT TCA GCA GAA CCT T-3'; sst5, forward primer, 5'-GGT TAC GTA GCA ACG ACC GTA CCA TCC GCA GCC CGG GTG-3'; reverse primer, 5'-GGT TCC GCC AGA GCT TC-3'. For 18 S rRNA evaluation, the TaqMan rRNA control reagents (Applied Biosystems; catalogue no. 4308329) were used according to the manufacturer's instructions. Data were analyzed using the relative standard-curve method, and results are expressed as relative expression normalized to 18 S rRNA expression.

Statistical analysis

Data were compared between groups using unpaired t-tests.

Results

Acute effect of SOM230 and octreotide

The levels of total and active ghrelin measured in fed animals were 2400±98 and 577±41 pg/ml, respectively. In order to increase the plasma levels of ghrelin, rats were fasted overnight. In fasted animals, the plasma concentrations of ghrelin reached the values of 3312±170 pg/ml for total ghrelin and 749±33 pg/ml for active ghrelin (Figs 1A and 2A). In both groups, the amount of active ghrelin represented about 20% of the amount of total ghrelin (24.0±1.0% in fed animals and 22.7±1.1% in fasted animals). In the treated groups, 1 h after s.c. injection of octreotide at 3, 10, or 30 μg/kg, the level of total ghrelin dropped down to the levels observed in fed animals (2451±86, 2293±120, and 2327±100 pg/ml, respectively; Fig. 1A). Stronger inhibition was observed on the level of active ghrelin, with levels below those observed in fed animals after treatment with octreotide 10 and 30 μg/kg (445±33 pg/ml, P = 0.02; 415±31 pg/ml, P = 0.008, respectively; Fig. 2A). When rats were fed ad libitum, a dose-dependent inhibition of total and active ghrelin secretion by octreotide was observed (Figs 1B and 2B), with levels significantly lower than in the non-treated group at doses of 10 and 30 μg/kg (1400±114 and 1030±133 pg/ml respectively for total ghrelin, and 371±28 and 274±27 pg/ml respectively for active ghrelin). SOM230 had no effect on total ghrelin at 3 or 30 μg/kg (Fig. 1) but, at 10 μg/kg, a small but significant increase in plasma ghrelin was observed in fasted rats (4148±166 versus 3312±170 pg/ml, P = 0.0032) and in fed rats (2249±122 versus 1770±108 pg/ml, P = 0.0093). Similarly, no effect of SOM230 on active ghrelin levels was observed at 3 μg/kg (Fig. 2). A tendency to further stimulate active ghrelin was observed at 10 μg/kg in fasted animals only. At 30 μg/kg, SOM230 inhibited active ghrelin secretion significantly in fasted rats.
but not in fed rats (Fig. 2B).

**Effects of prolonged treatment with SOM230 and octreotide**

Rats treated for 14 days with SOM230 or octreotide had a body weight lower than rats receiving 0.9% NaCl, thus confirming previous data (26) and the efficiency of the compound infusion (Fig. 3). After 7 and 14 days of treatment, octreotide significantly inhibited total ghrelin increase induced by overnight fasting, as well as total ghrelin levels in fed rats (Fig. 4). In contrast, SOM230 had no effect on total ghrelin levels after 7 and 14 days of treatment at any dose. After 7 days of treatment, fasting-induced active ghrelin levels were inhibited strongly (below levels of fed rats) by both octreotide and SOM230, with a stronger effect observed with octreotide (Fig. 5A, left-hand panel). After 14 days, loss of response was observed for both compounds. Complete loss of inhibitory effect was observed for SOM230, and partial loss for octreotide, with levels of active ghrelin still reduced to the levels observed (Fig. 5A, right-hand panel). In fed rats, after 7 days, octreotide inhibited active ghrelin secretion (647 ± 54 pg/ml for vehicle-treated rats, and 283 ± 67 pg/ml, P = 0.001; 482 ± 70 pg/ml, P = 0.07; 359 ± 44 pg/ml, P = 0.0012 at 3, 10, and 30 μg/kg per h, respectively), and SOM230 also inhibited active ghrelin secretion at 30 μg/kg per h (440 ± 47 pg/ml, P = 0.01). After 14 days, the inhibitory effect of both compounds was reduced (Fig. 5B).

The expression of somatostatin precursor mRNA and sst receptor mRNAs was measured by real-time reverse transcriptase PCR in biopsies of glandular stomach from fed or overnight fasted rats. A predominant expression of sst2 and sst1 was measured in the rat glandular stomach. sst3 and sst4 mRNAs were also expressed in

---

**Figure 2** Acute effect of SOM230 and octreotide on (A) fasting-induced active ghrelin secretion and (B) active ghrelin secretion in fed rats. Rat active ghrelin plasma levels as determined by RIA are expressed as mean ± S.E.M.; n = 6 animals per group. Rats were fed ad libitum or fasted overnight, and SOM230 and octreotide were injected s.c. 1 h before blood sampling. **P < 0.01, ***P < 0.001 compared with the fed group. §P < 0.05, §§P < 0.01, §§§P < 0.001 compared with the fasted group.

---

**Figure 3** Change in rat body weight 7 and 14 days after s.c. implantation of osmotic minipumps (MP). Rats were either fasted (A) or fed ad libitum (B). Results are expressed as mean ± S.E.M. of the percentage of initial body weight; n = 6 rats per group. • with solid line, fed rats; ● with dashed line, fasted rats; ● with dashed line, octreotide 3 μg/kg per h; ▲ with dashed line, octreotide 10 μg/kg per h; ■ with dashed line, octreotide 30 μg/kg per h; ● with long/short-dashed line, SOM230 3 μg/kg per h; ▲ with long/short-dashed line, SOM230 10 μg/kg per h; ■ with long/short-dashed line, SOM230 30 μg/kg per h.
lower amounts, and ssr5 mRNA was barely detectable in the rat glandular stomach (Fig. 6). ssr2 receptor expression was lower in fasted compared with fed animals (relative expression of $2.23^{0.09}\text{pg/ml in fed rats compared to }1.13^{0.33}\text{pg/ml in fasted rats, }P=0.018$), in contrast to ssr1, ssr3, and ssr4. Somatostatin precursor mRNA was also expressed in significant amounts in the rat glandular stomach. Its expression was not affected by overnight fasting.

**Discussion**

This study shows the dose-dependent inhibition of ghrelin secretion achieved 1 h following injection of octreotide at 3, 10, and 30 $\mu$g/kg, both in fed rats and after stimulation of secretion by overnight fasting. With SOM230, significant acute inhibition of active ghrelin was observed at 30 $\mu$g/kg in fasted rats only. After 7 days of treatment, active ghrelin was strongly inhibited by both octreotide and SOM230 in fasted rats, with a stronger effect for octreotide. This inhibition of active ghrelin secretion after 7 days was less pronounced in fed rats. After 14 days, desensitization of this effect was observed, leading to lower inhibition with octreotide and absence of inhibition with SOM230 in fasted rats, and absence of inhibition for both compounds in fed rats. Since food intake is stimulated by the octanoylated form of ghrelin (2), those data obtained for active ghrelin modulation are the more physiologically relevant. ssr receptor expression analysis in the rat glandular stomach showed a predominant expression of ssr1 and ssr2, lower ssr3 and ssr4, and barely detectable ssr5 mRNA expression. These data suggest that inhibition of ghrelin secretion might be mediated by ssr1, ssr2, or ssr3 receptors. The involvement of ssr4 and ssr5 receptors can be excluded because both compounds tested have no activity at the ssr4 receptor, and ssr5 receptor is not expressed in the target tissue. The affinity of octreotide for ssr2 receptors is about 3-fold higher compared with SOM230, with IC50 values of 0.38 and 1.0 nM, respectively (25). In contrast, the affinity of SOM230 for ssr1 and ssr3 receptors is higher compared with octreotide (9.3 and 1.5 nM versus 280 and 7.1 nM, respectively (25)), thus excluding ssr1 and ssr3 as predominant mediators of the inhibition of ghrelin secretion in rats. Recent functional data obtained with cells expressing human recombinant ssr receptors showed a 6-fold higher functional activity of octreotide compared with SOM230 on ssr2 receptors in terms of inhibition of forskolin-induced cAMP.
This difference of efficacy is in line with the differences observed for the inhibition of active ghrelin secretion by octreotide and SOM230. Therefore, the sst2 receptor is the most relevant mediator of the inhibition of ghrelin secretion by octreotide and SOM230 in rats.

After 14 days of continuous treatment the inhibitory effect of SOM230 and octreotide declined. Desensitization was previously observed for the regulation of GH/IGF-I secretion by octreotide (26). Inhibition of the GH/IGF-I axis by somatostatin is a sst2-/sst5-mediated effect (28), and stronger desensitization was observed with the sst2-prefering analogue octreotide compared with the multireceptor ligand SOM230 (26). Therefore, the similarity of the desensitization observed with octreotide and SOM230 on active ghrelin secretion and the difference observed on GH and IGF-1 secretion further argue for a predominantly or exclusively sst2-mediated regulation of ghrelin plasma levels in the rat.

Figure 5 Effect of SOM230 and octreotide on active ghrelin secretion after 7 and 14 days of treatment. Rats were either (A) fasted overnight or (B) fed ad libitum. Rat active ghrelin plasma levels as determined by RIA on days 7 (left-hand panels) and 14 (right-hand panels) are expressed as mean±S.E.M.; n=6 animals per group. *P<0.05, **P<0.01 compared with the fed group. §§P<0.05, §§§P<0.01 compared with the fasted group.

Figure 6 Expression of somatostatin (SRIF) and sst receptor subtype mRNAs in the glandular stomach of overnight fasted rats (left-hand panel) and fed rats (right-hand panel) was determined by quantitative reverse transcriptase PCR. Results are expressed as mean±S.E.M. from target mRNA expression normalized to 18S rRNA expression, in arbitrary units (A.U.); n=5 rats per group.
Somatostatin receptor expression analysis showed downregulation of sst₂ receptors in the rat glandular stomach after overnight fasting. Although the present data do not show an effect of fasting on somatostatin precursor mRNA expression, the downregulation of sst₂ mRNA expression might be explained by an increased somatostatin peptide content in the rat glandular stomach. Indeed, various studies have shown that somatostatin concentration in the antral stomach of the rat is increased after fasting (29–31). In a study by Roca et al. (31), increased somatostatin concentration was associated with a decreased number of somatostatin binding sites in the antral mucosa.

In the acute-treatment experiments, after 1 h, a further increase in total ghrelin plasma levels (and a trend to increase in active ghrelin) was observed in the groups treated with 10 μg/kg SOM230. At 30 μg/kg SOM230, the expected inhibition of active ghrelin was observed in fasted rats. The unexpected increase in ghrelin observed with 10 μg/kg SOM230 might be explained by the difference in binding affinities between SOM230 and octreotide on sst₂ and sst₅ receptors, resulting in different effects on hormone secretion. In binding experiments octreotide has a 16-fold higher affinity for sst₂ compared with sst₅ receptors. In contrast, SOM230 has a 6-fold higher affinity for sst₅ compared with sst₂ receptors. According to our study, sst₂ receptor is hardly expressed in the stomach, i.e. the main source of ghrelin secretion, in contrast to insulin secreting β-cells of the pancreas (32). The predominant expression of sst₅ receptors on β-cells is the histological correlate for the 11-fold stronger inhibitory effect of SOM230 on insulin secretion compared with octreotide (25). Several studies have shown that circulating insulin has an inhibitory effect on ghrelin secretion (33–35). Therefore, by inhibiting insulin secretion via the activation of sst₅ receptors in the pancreas, SOM230 at 10 μg/kg could disinhibit the inhibitory effect of insulin on ghrelin secretion, leading to a further increase in ghrelin. At 30 μg/kg the concentration of SOM230 could be high enough to directly inhibit ghrelin secretion from the stomach via the stimulation of sst₂ receptors, and thereby overcome the indirect effect caused by the sst₅-mediated inhibition of insulin. The stronger inhibitory effect of octreotide on ghrelin secretion can be explained by the 6-fold-higher affinity for and the 6-fold-higher functional activity at sst₂ receptors compared with SOM230 (27).

Our data are in line with previously published data showing that somatostatin and octreotide at nanomolar concentrations inhibited total ghrelin secretion from the perfused rat stomach (36). The inhibition of ghrelin secretion by octreotide has also been shown in a study on nine young healthy male volunteers, with plasma ghrelin concentrations representing almost 50% of the values obtained under saline infusion (37). In acromegalic patients treated with octreotide, a significant decrease in plasma ghrelin levels was observed (20). The decrease in human plasma levels of ghrelin was observed after 3–9 months of treatment with octreotide. This suggests that, in humans, the inhibitory effect of octreotide on ghrelin secretion is long lasting with slower desensitization than that observed in our study in the rat. A more recent study investigating the effect of somatostatin i.v. infusion in PWS patients showed that somatostatin can also reduce the levels of plasma ghrelin in these patients, although it did not change food intake measured as the quantity of food taken during a period of 1 h (38).

In summary, the present study indicates that somatostatin analogues may be useful for the inhibition of stimulated and unstimulated ghrelin plasma levels. In the rat, this appears to be mediated primarily or exclusively by the sst₂ receptor. In rats rapid desensitization is seen after 14 days, in contrast to the observations made by others in humans. These observations suggest that other somatostatin receptors besides sst₂ might be more relevant for the inhibitory effect on ghrelin secretion in humans compared with rats, with potential implications for multireceptor ligands such as SOM230.

Acknowledgements

We are thankful to Ramona Rebmann for her skillful technical assistance. We also would like to thank Dr Christian Bruns and Professor Gisbert Weckbecker for their support, as well as Gérard Vogt, Cyril Allard, and Corinne Delucis-Bronn for their technical advice.

References


